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NiceProt View of Swiss- Prot: Q64280

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Entry information

Entry name	TGF4_MOUSE
Primary accession number	Q64280
Secondary accession numbers	None
Entered in Swiss-Prot in	Release 35, November 1997
Sequence was last modified in	Release 35, November 1997
Annotations were last modified in	Release 44, July 2004
Name and origin of the protein	
Protein name	Transforming growth factor beta 4 [Precursor]
Synonyms	TGF-beta 4 Lefty protein Lefty-1 protein STR43 protein
Gene name	Name: Ebaf Synonyms: Tgfb4, Stra3, Lefty, Lefty1
From	Mus musculus (Mouse) [TaxID: 10090]
Taxonomy	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

References

[1] SEQUENCE FROM NUCLEIC ACID.
 DOI=[10.1038/381151a0](https://doi.org/10.1038/381151a0); MEDLINE=96202359; PubMed=8610011 [NCBI, ExPASy, EBI, Israel, Japan]
Meno C., Saijoh Y., Fujii H., Ikeda M., Yokoyama T., Yokoyama M., Toyoda Y., Hamada H.;
 "Left-right asymmetric expression of the TGF beta-family member lefty in mouse embryos.";
 Nature 381:151-155(1996).

[2] SEQUENCE FROM NUCLEIC ACID.
Bouillet P.;
 Submitted (JUN-1996) to the EMBL/GenBank/DDBJ databases.

[3] SEQUENCE FROM NUCLEIC ACID.

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<h1>NiceProt View of Swiss- Prot: O00292</h1>				
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Note: most headings are clickable, even if they don't appear as links. They link to the user manual or other documents.

Entry information

Entry name **TGF4_HUMAN**
 Primary accession number **O00292**
 Secondary accession numbers **O75611 Q8NBQ9**
 Entered in Swiss-Prot in **Release 35, November 1997**
 Sequence was last modified in **Release 40, October 2001**
 Annotations were last modified in **Release 44, July 2004**

Name and origin of the protein

Protein name **Transforming growth factor beta 4 [Precursor]**
 Synonyms **TGF-beta 4**
Endometrial bleeding-associated factor
Left-right determination factor A
Lefty-A protein

Gene name

Name: EBAF

Synonyms: **TGFB4, LEFTA, LEFTYA**

Homo sapiens (Human) [TaxID: 9606]

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

References

[1] SEQUENCE FROM NUCLEIC ACID.

TISSUE=Placenta;

MEDLINE=97298127;PubMed=9153275 [NCBI, ExPASy, EBI, Israel, Japan]

Kothapalli R., Buyukal I., Wu S.-Q., Chegini N., Tabibzadeh S.;

"Detection of ebaf, a novel human gene of the transforming growth factor beta superfamily association of gene expression with endometrial bleeding.";
J. Clin. Invest. 99:2342-2350(1997).

[2] REVISIONS.

MEDLINE=99162193;PubMed=10053005 [NCBI, ExPASy, EBI, Israel, Japan]

Kothapalli R.;

Unpublished results, cited by: Kosaki K., Bassi M.T., Kosaki R., Lewin M., Belmont J., Schauer G.,

Casey B.; Am. J. Hum. Genet. 64:712-721(1999).

[3] SEQUENCE FROM NUCLEIC ACID, AND VARIANT L-R AXIS MALFORMATIONS ASN-342.

TISSUE=Placenta;

MEDLINE=99162193;PubMed=10053005 [NCBI, ExPASy, EBI, Israel, Japan]

Kosaki K., Bassi M.T., Kosaki R., Lewin M., Belmont J., Schauer G., Casey B.;

"Characterization and mutation analysis of human LEFTY A and LEFTY B, homologues of murine genes implicated in left-right axis development.";

Am. J. Hum. Genet. 64:712-721(1999).

[4] SEQUENCE FROM NUCLEIC ACID.

DOI=10.1038/ng1285;PubMed=14702039 [NCBI, ExPASy, EBI, Israel, Japan]

Ota T., Suzuki Y., Nishikawa T., Otsuki T., Sugiyama T., Irie R., Wakamatsu A., Hayashi K., Sato H., Nagai K., Kimura K., Makita H., Sekine M., Obayashi M., Nishi T., Shibahara T., Tanaka T., Ishii S., Yamamoto J.-I., Saito K., Kawai Y., Isono Y., Nakamura Y., Nagahari K., Murakami K., Yasuda T., Iwayanagi T., Wagatsuma M., Shiratori A., Sudo H., Hosoiri T., Kaku Y., Kodaira H., Kondo H., Sugawara M., Takahashi M., Kanda K., Yokoi T., Furuya T., Kikkawa E., Omura Y., Abe K., Kamihara K., Katsuta N., Sato K., Tanikawa M., Yamazaki M., Ninomiya K., Ishibashi T., Yamashita H., Murakawa K., Fujimori K., Tanai H., Kimata M., Watanabe M., Hiraoka S., Chiba Y., Ishida S., Ono Y., Takiguchi S., Watanabe S., Yosida M., Hotuta T., Kusano J., Kanehori K., Takahashi-Fujii A., Hara H., Tanase T.-O., Nomura Y., Togoya S., Komai F., Hara R., Takeuchi K., Arita M., Imose N., Musashino K., Yuuki H., Oshima A., Sasaki N., Aotsuka S., Yoshikawa Y., Matsunawa H., Ichihara T., Shiohata N., Sano S., Moriya S., Momiyama H., Satoh N., Takami S., Terashima Y., Suzuki O., Nakagawa S., Senoh A., Mizoguchi H., Goto Y., Shimizu F., Wakebe H., Hishigaki H., Watanabe T., Sugiyama A., Takemoto M., Kawakami B., Yamazaki M., Watanabe K., Kumagai A., Itakura S., Fukuzumi Y., Fujimori Y., Komiyama M., Tashiro H., Tanigami A., Fujiwara T., Ono T., Yamada K., Fujii Y., Ozaki K., Hirao M., Ohmori Y., Kawabata A., Hikiji T., Kobatake N., Inagaki H., Ikema Y., Okamoto S., Okitani R., Kawakami T., Noguchi S., Itoh T., Shigeta K., Senba T., Matsumura K., Nakajima Y., Mizuno T., Morinaga M., Sasaki M., Togashi T., Oyama M., Hata H., Watanabe M., Komatsu T., Mizushima-Sugano J., Satoh T., Shirai Y., Takahashi Y., Nakagawa K., Okumura K., Nagase T., Nomura N., Kikuchi H., Masuho Y., Yamashita R., Nakai K., Yada T., Nakamura Y., Ohara O., Isogai T., Sugano S.;

"Complete sequencing and characterization of 21,243 full-length human cDNAs.";

Nat. Genet. 36:40-45(2004).

[5] SEQUENCE FROM NUCLEIC ACID.

TISSUE=Ovary;

DOI=10.1073/pnas.242603899;MEDLINE=22388257;PubMed=12477932 [NCBI, ExPASy, EBI, Israel, Japan]

Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G., Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D., Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K., Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F., Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L., Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E., Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C., Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullahy S.J., Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H., Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W., Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A., Fahey J., Helton E., Ketteman M., Madan A., Rodrigues S., Sanchez A., Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G., Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C., Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M., Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smailus D.E., Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;

"Generation and initial analysis of more than 15,000 full-length human and mouse cDNA

sequences.";
Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).

Comments

- **FUNCTION:** Required for left-right (L-R) asymmetry determination of organ systems in mammals. May play a role in endometrial bleeding.
- **SUBCELLULAR LOCATION:** Secreted.
- **TISSUE SPECIFICITY:** Mesenchymal cells of the endometrial stroma.
- **DEVELOPMENTAL STAGE:** Transiently expressed before and during menstrual bleeding.
- **PTM:** The processing of the protein may also occur at the second R-X-X-R site located at AA 132-135. Processing appears to be regulated in a cell-type specific manner.
- **DISEASE:** Defects in EBAF are the cause of left-right axis malformations (L-R axis malformation) [MIM:601877]. The defect includes left pulmonary isomerism, with cardiac anomalies characterized by complete atrioventricular canal defect and hypoplastic left ventricle, and interrupted inferior vena cava.
- **SIMILARITY:** Belongs to the TGF-beta family.

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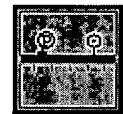
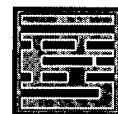
Cross-references

	U81523; AAB53269.1; ALT_SEQ. [EMBL / GenBank / DDBJ] [CoDingSequence]
	AF081511; AAC32600.1; -. [EMBL / GenBank / DDBJ] [CoDingSequence]
	AF081508; AAC32600.1; JOINED. [EMBL / GenBank / DDBJ] [CoDingSequence]
EMBL	AF081509; AAC32600.1; JOINED. [EMBL / GenBank / DDBJ] [CoDingSequence]
	AF081510; AAC32600.1; JOINED. [EMBL / GenBank / DDBJ] [CoDingSequence]
	AF081513; AAD48145.1; -. [EMBL / GenBank / DDBJ] [CoDingSequence]
	AK075344; BAC11556.1; -. [EMBL / GenBank / DDBJ] [CoDingSequence]
	BC035718; AAH35718.1; -. [EMBL / GenBank / DDBJ] [CoDingSequence]
HSSP	P10600 ; 1TGJ. [HSSP ENTRY / PDB]
Genew	HGNC:3122 ; EBAF.
CleanEx	HGNC:3122 ; EBAF.
GeneCards	EBAF .
GeneLynx	EBAF ; <i>Homo sapiens</i> .
GenAtlas	EBAF .
MIM	601877 [NCBI / EBI]. GO:0007275 ; Biological process: development (<i>traceable author statement</i>). GO:0007309 ; Biological process: oocyte axis determination (<i>traceable author statement</i>).
GO	GO:0007179 ; Biological process: transforming growth factor beta receptor signaling pathway (<i>traceable author statement</i>). QuickGo view .
SOURCE	EBAF ; <i>Homo sapiens</i> .
Ensembl	O00292; <i>Homo sapiens</i> . [Entry / Contig view] IPR001839 ; TGFb.
InterPro	IPR001111 ; TGFb_N. Graphical view of domain structure .

Pfam	PF00688 ; TGFb_propeptide; 1. PF00019 ; TGF_beta; 1. Pfam graphical view of domain structure .
ProDom	PD000357 ; TGFb; 1. [Domain structure / List of seq. sharing at least 1 domain]
PROSITE	PS00250 ; TGF_BETA_1; 1.
HOVERGEN	[Family / Alignment / Tree]
BLOCKS	Q00292 .
ProtoNet	Q00292 .
ProtoMap	Q00292 .
PRESAGE	Q00292 .
DIP	Q00292 .
ModBase	Q00292 .
SMR	Q00292 ; 63A416CAE30F7A39.
SWISS-2DPAGE	Get region on 2D PAGE .
UniRef	View cluster of proteins with at least 50% / 90% identity.

Keywords

[Cytokine](#); [Developmental protein](#); [Disease mutation](#); [Glycoprotein](#); [Growth factor](#); [Multigene family](#); [Signal](#).

Features[Feature table viewer](#)[Feature aligner](#)

Key	From	To	Length	Description	FTId
SIGNAL	1	21	21	Potential.	
PROPEP	22	76	55	Or 135 (Potential).	
CHAIN	77	366	290	Transforming growth factor beta 4.	
DISULFID	251	264	13	By similarity.	
DISULFID	263	316	53	By similarity.	
DISULFID	293	351	58	By similarity.	
DISULFID	297	353	56	By similarity.	
CARBOHYD	158	158	0	N-linked (GlcNAc...) (Potential).	
VARIANT	342	342	0	*	S -> N (in L-R axis malformations). VAR_010385
CONFLICT	183	183	0	A -> P (in Ref. 4).	

Sequence information

Length: 366 AA [This is the length of the unprocessed precursor]

Molecular weight: 40920 Da [This is the MW of the unprocessed precursor]

CRC64: 63A416CAE30F7A39 [This is a checksum on the sequence]

10	20	30	40	50	60
MWPLWL	WVLPLAGPGA	ALTEEQ	LLRQLQLSEV	PVLDRADMEK	LVIPAHVRAQ
RRSHG	DRSRGKRF	SFREVAGRFL	ASEASTHLLV	FGMEQRLPPN	SELVQAVLRL
70	80	90	100	110	120

130 FQEPVPKAAL 140 HRHGRLSPRS 150 AQARVTVEWL 160 RVRDDGSNRT 170 SLIDSRLVSV 180 HESGWKAFDV
 190 TEAVNFWQQL 200 SRPRQPLLQ 210 VSVQREHLGP 220 LASGAHKLVR 230 FASQGAPAGL 240 GEPQLELHTL
 250 DLRDYGAQGD 260 CDPEAPMTEG 270 TRCCRQEMYI 280 DLQGMKWAKN 290 WVLEPPGFLA 300 YECVGTCCQP
 310 PEALAFNWPFF 320 LGPRQCIASE 330 TASLPMIVSI 340 KEGGRTRPQV 350 VSLPNMRVQK 360 CSCASDGALV
 PRRLQP

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Sequence analysis tools: [ProtParam](#), [ProtScale](#),
[Compute pI/Mw](#), [PeptideMass](#), [PeptideCutter](#),
[Dotlet](#) (Java)



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Search in Swiss-Prot and TrEMBL for: tgfb4

Swiss-Prot Release 44.5 of 13-Sep-2004

TrEMBL Release 27.5 of 13-Sep-2004

- Number of sequences found in Swiss-Prot₍₂₎ and TrEMBL₍₀₎: 2
- For more directed searches, you can use the Sequence Retrieval System SRS.

Search in Swiss-Prot: There are matches to 2 out of 158316 entries

TGF4_HUMAN (O00292)

Transforming growth factor beta 4 precursor (TGF-beta 4) (Endometrial bleeding-associated factor) (Left-right determination factor A) (Lefty-A protein). {GENE: Name=EBAF; Synonyms=TGFB4, LEFTA, LEFTYA} - Homo sapiens (Human)

TGF4_MOUSE (Q64280)

Transforming growth factor beta 4 precursor (TGF-beta 4) (Lefty protein) (Lefty-1 protein) (STRA3 protein). {GENE: Name=Ebaf; Synonyms=Tgfb4, Stra3, Lefty, Lefty1} - Mus musculus (Mouse)

Search in TrEMBL: There are matches to 0 out of 1400820 entries

in Swiss-Prot/TrEMBL by AC, ID, description,
gene name, organism

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7/3, KWIC/5 (Item 1 from file: 399)

DIALOG(R) File 399:CA SEARCH(R)

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111091468 CA: 111(11)91468h JOURNAL

Complementary deoxyribonucleic acid cloning of a messenger ribonucleic acid encoding transforming growth factor β 4 from chicken embryo chondrocytes

AUTHOR(S): Jakowlew, Sonia B.; Dillard, Pamela J.; Sporn, Michael B.; Roberts, Anita B.

LOCATION: Lab. Chemoprev., Natl. Cancer Inst., Bethesda, MD, 20892, USA

JOURNAL: Mol. Endocrinol. DATE: 1988 VOLUME: 2 NUMBER: 12 PAGES: 1186-95 CODEN: MOENEN ISSN: 0888-8809 LANGUAGE: English

? t s17/9/3 4

17/9/3 (Item 1 from file: 73)

DIALOG(R) File 73:EMBASE

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06240683 EMBASE No: 1995269574

The immunomodulatory diversity of the proteins of the transforming growth factor β (TGF β P) family

Wieczorek Z.; Sion J.; Kluczyk A.; Zbozien R.; Stafanowicz P.; Siemion I.Z.

L. Hirschfeld Inst Immun/Exp Therapy, Polish Academy of Sciences, Czerska 12, 53-114 Wroclaw Poland

International Journal of Peptide and Protein Research (INT. J. PEPT.

PROTEIN RES.) (Denmark) 1995, 46/2 (113-118)

CODEN: IJPPC ISSN: 0367-8377

DOCUMENT TYPE: Journal; Article

LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH

The examination of immunomodulatory properties of oligopeptides derived from two exposed loops (containing thymopentin-like and tuftsin-like sequences, respectively) of the proteins belonging to TGF β family suggests that the particular species of the TGF β family should differ distinctly in their influence on the immune response. According to our results obtained from three TGF β species of mammals, TGF β 2 should be a strong immunosuppressor, whereas for TGF β 3 the immunostimulative potency is more probable. TGF β 1 species would possess both immunosuppressive and immunostimulative potency, residing in two different loops of the protein. The results obtained also suggest that chicken TGF β 4 should be associated with immunostimulative effects and xenopus TGF β 5 with immunosuppressive ones.

DRUG DESCRIPTORS:

*transforming growth factor beta--pharmacology--pd; *transforming growth factor beta--drug comparison--cm; *transforming growth factor beta--drug analysis--an; *transforming growth factor beta--drug development--dv thymopentin--drug comparison--cm

MEDICAL DESCRIPTORS:

*delayed hypersensitivity; *immunomodulation animal experiment; article; controlled study; drug purification; drug synthesis; high performance liquid chromatography; mouse; nonhuman; protein variant

CAS REGISTRY NO.: 69558-55-0 (thymopentin)

SECTION HEADINGS:

026 Immunology, Serology and Transplantation

029 Clinical and Experimental Biochemistry

030 Clinical and Experimental Pharmacology

037 Drug Literature Index

17/9/4 (Item 2 from file: 73)

DIALOG(R) File 73:EMBASE

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06190248 EMBASE No: 1995227311

Expression of transforming growth factor beta in the embryonic avian lens coincides with the presence of mitochondria

Potts J.D.; Bassnett S.; Beebe D.C.

Department of Anatomy/Cell Biology, Uniformed Svcs. Univ. of Health Sci.,
4301 Jones Bridge Road, Bethesda, MD 20814-4799 United States

Developmental Dynamics (DEV. DYN.) (United States) 1995, 203/3
(317-323)

CODEN: DEDYE ISSN: 1058-8388

DOCUMENT TYPE: Journal; Article

LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH

During their maturation, lens cells lose all membrane bound organelles, including mitochondria. In chicken embryos this process begins in the central lens fibers beginning around embryonic day 12 (E12). Transforming growth factor beta (TGF β) is a multipotent growth modulator thought to play a role in numerous developmental processes. TGF β 1 has been localized to mitochondria in rat liver cells and muscle cells. In the present study, we examined the expression of TGF β isoform mRNAs and proteins during chicken embryonic lens development. PCR analysis demonstrated TGF β 2 and TGF β 3 transcripts in the lens epithelium and fibers throughout pre- and post-hatching development. TGF β isoforms were detected throughout the lens epithelium and fibers early in development (E6). However by E19, the distribution of TGF β 2 and TGF β 3 transcripts and proteins coincided with regions of the lens that contained mitochondria. In addition, intense TGF β staining was observed in the basal portions of the equatorial epithelial cells, a region with abundant mitochondria. Transcripts for TGF β 1 and TGF β 4 were not detected in any tissue or time frame examined. Similarly, no immunostaining for TGF β 1 was observed.

DRUG DESCRIPTORS:

*transforming growth factor beta

MEDICAL DESCRIPTORS:

*embryo development; *gene expression; *lens animal cell; article; cell differentiation; cell maturation; cell organelle; chick embryo; embryo; immunoblotting; immunohistochemistry; lens epithelium; mitochondrion; nonhuman; polymerase chain reaction; priority journal

SECTION HEADINGS:

021 Developmental Biology and Teratology

Cloning and characterization of human polyamine-modulated factor-1, a transcriptional cofactor that regulates the transcription of the spermidine/spermamine N(1)-acetyltransferase gene.

Wang Y; Devereux W; Stewart T M; Casero R A

Johns Hopkins Oncology Center Research Laboratories, Baltimore, Maryland 21231, USA.

Journal of biological chemistry (UNITED STATES) Jul 30 1999, 274 (31)

p22095-101, ISSN 0021-9258 Journal Code: 2985121R

Contract/Grant No.: CA51085; CA; NCI; CA58184; CA; NCI

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: Completed

Subfile: INDEX MEDICUS

The increased transcription and ultimate superinduction of the spermidine/spermamine N(1)-acetyltransferase (SSAT) gene has been associated with the antineoplastic activity of several new antitumor polyamine analogues. In sensitive tumor cell types, the transcriptional induction appears to be regulated by the constitutive association of the transcription factor Nrf-2 with the recently discovered polyamine-responsive element. Using the yeast two-hybrid system, a new transcriptional cofactor, polyamine-modulated factor-1 (PMF-1), has been identified as a partner protein of Nrf-2 that, in combination with Nrf-2, regulates the polyamine analogue-induced transcription of SSAT. The human

PMF - 1 gene, located on chromosome 1 near the 1q12/1q21 border, yields an mRNA transcript of approximately 1.2 kilobases that codes for a 165-amino acid protein with a predicted molecular mass of approximately 20 kDa. The PMF-1 mRNA appears to increase in response to analogue exposure only in analogue-responsive cells. In addition to the transcriptional regulation of SSAT, PMF-1 or similar factors should be considered in the regulation of other polyamine-dependent genes.

Tags: Female; Human; Pregnancy; Support, U.S. Gov't, P.H.S.

Descriptors: *Acetyltransferases--genetics--GE; *Chromosomes, Human, Pair 1; *Gene Expression Regulation, Enzymologic; *Transcription Factors --genetics--GE; *Transcription Factors--metabolism--ME; *Transcription, Genetic; Amino Acid Sequence; Base Sequence; Cell Line; Chromosome Mapping; Cloning, Molecular; DNA Primers; DNA-Binding Proteins--metabolism--ME; Exons; Gene Library; Introns; Molecular Sequence Data; Molecular Weight; Placenta--metabolism--ME; RNA, Messenger--genetics--GE; Recombinant Proteins--chemistry--CH; Transcription Factors--chemistry--CH; Transfection Molecular Sequence Databank No.: GENBANK/AF141308; GENBANK/AF141309; GENBANK/AF141310; GENBANK/AH008078

CAS Registry No.: 0 (DNA Primers); 0 (DNA-Binding Proteins); 0 (RNA, Messenger); 0 (Recombinant Proteins); 0 (Transcription Factors); 0 (nuclear respiratory factor 2); 0 (polyamine-modulated factor 1)

Enzyme No.: EC 2.3.1. (Acetyltransferases); EC 2.3.1.57 (spermidine acetyltransferase)

Record Date Created: 19990819

Record Date Completed: 19990819